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(54) Aqueous Wellbore Service Fluids

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AQUEOUS WELLBORE SERVICE FLUIDS

This invention relates to aqueous wellbore service fluids, including drilling fluids, completion fluids, work over fluids, packer fluids, fracturing fluids and the like, which may be employed in various well servicing operations. More specifically, it relates to thickened, substantially solids-free high electrolyte-containing aqueous fluids which are employed as a base fluid to prepare many types of wellbore service fluids.

Essentially solids-free aqueous fluids containing electrolytes have some advantages over clay-based fluids for preparing wellbore service fluids because: (a) they do not normally contain undesirable solids which can cause formation damage, (b) they contain hydration inhibiting materials such as potassium chloride, calcium chloride or the like, which are important to prevent damage to clay containing formations, and (c) they can be prepared over a wide range of densities.

The viscosity of high electrolyte-containing aqueous fluids is, however, difficult to control because of the high electrolyte concentration. Thickened fluids

are desirable for carrying solids, e.g., in cleaning out wells, drilling and the like. Likewise, thickened fluids resist water loss, which may be damaging to petroleum producing subterranean formations.

5 Bydroxy alkyl celluloses have been employed to thicken electrolyte-containing aqueous fluids to improve the solid carrying capacity thereof. Likewise, starch has been employed to aid in water loss control of these fluids, but with limited success. However, 10 these materials are difficult to disperse and dissolve in concentrated electrolytes at ambient temperature; the viscosity of the resulting solutions tend to decrease with an increase in temperature; and the hydroxy alkyl celluloses are subject to shear degradation under normal operating conditions.

Certain quaternary ammonium salts have been shown to impart viscoelastic properties to aqueous solutions, S. Gravsholt "Viscoelasticity in Highly Dilute Aqueous Solutions of Puze Cationic Detergents", Journal of Colloid and Interface Science, Vol.57, No. 20 3, December 1976, pp. 575-577. Gravsholt showed that cetyl trimethyl ammonium bromide would not impart viscoelastic properties to water but that cetyl trimethyl ammonium salicylate and certain other aromatic 25 anion-containing quaternary amines would. In U.S. Patent 3,292,698, a mixture of cyclohexyl ammonium chloride and undecane-3-sodium sulfate was taught to induce viscoelastic properties to a formation flooding liquid containing less than about 3.5 percent by weight of sodium chloride. Higher levels of sodium chloride 30 were said to destroy the viscoelastic properties of the fluid. In British Patent No. 1,443,244, a specific

ethoxylated or propoxylated tertiary amine is employed to thicken an aqueous solution of a strong mineral acid. U.S. Patent 3,917,536 teaches that certain primary amines may be employed in subterranean formation acidizing solutions to retard the reaction of the acid on the formation. The amine may be more readily dispersed into the acid solution with the use of a dispersing agent such as a quaternary amine.

It is a feature of the present invention to provide a high electrolyte-containing aqueous wellbore service fluid which has improved viscosity characteristics over a wide range of wellbore conditions; is easier to prepare at the well site and has better shear stability and consistent viscosity over a wide temperature range.

The improved aqueous wellbore service fluid of the present invention can be employed in well-known wellbore services such as, perforation, clean-up, long term shut-ins, drilling, placement of gravel packs, and the like. These services are well known in the art and are taught, for example, in U. S. Patent Nos. 3,993,570; 3,176,950; 3,126,950; 2,898,294 and in C. M. Hudgens et al "High Density Packer Fluids Pay Off in South Louisiana": World Oil, 1961, pp. 113-119.

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As employed herein, "ppg" means pounds per gallon. Also, when "percent" or "%" are employed, they mean percent by weight unless otherwise specified.

The present invention comprises an aqueous wellbore service fluid comprising:

water, a sufficient quantity of at least one water soluble salt to increase the density of said fluid to within a range of from 12 to 21 lbs/gal, and a sufficient quantity of at least one thickener soluble in said fluid to increase the viscosity of said fluid to at least 50% over the viscosity of the salt containing fluid, said thickener being at least one member selected from the group consisting of: (a) an amine corresponding to the formula

$$R_1 - N$$
 wherein R_3

R₁ is at least about a C₁₆ aliphatic group which may be branched or straight chained and which may be saturated or unsaturated;

 R_2 and R_3 are each independently, hydrogen or a C_1 to about C_6 aliphatic group which can be branched or straight chained, saturated or unsaturated and which may be substituted with a group which renders the R_2 and/or R_3 group more hydrophilic; (b) salts of said amine corresponding to the formula

$$R_1 - N^+ - H X^-$$
 wherein

 $\rm R_1,\ R_2$ and $\rm R_3$ are the same as defined hereinbefore and X is an inorganic or organic salt forming anion; or (c) a quaternary ammonium salt of said amine corresponding to the formula

$$R_1 - N^+ - R_4 X^-$$
 wherein R_3

 R_1 , R_2 , R_3 and X^- are the same as hereinbefore defined and R_4 independently constitutes a group which has previously been set forth for R_2 and R_3 , none of R_1 , R_2 , R_3 or R_4 are hydrogen, and the R_2 , R_3 and R_4 groups of the amine salt and quaternary ammonium salt may be formed into a heterocyclic 5 or 6 member ring structure which includes the nitrogen atom of the amine.

The aqueous wellbore service fluid may have a density ranging from as low as about 8.5 ppg, preferably about 12 ppg, to about 21 ppg. It has been found that the higher density fluids are more difficult to thicken because of the high electrolyte content. It is at these higher densities, e.g. about 15 ppg and higher, that the practice of the present invention is particularly useful. However, advantages are also achieved in the lower density fluids.

The density is achieved by dissolving one or more water soluble inorganic salts in water to provide a substantially solids-free fluid. Naturally occurring brines and seawater can be employed if desired. Preferably, the aqueous wellbore service fluid contains at

least about 3 percent of a water soluble salt of potassium. calcium or sodium. In addition, the aqueous fluid may contain other soluble salts of, for example, zinc, lithium, chromium, iron, copper, and the like. Preferably inorganic chlorides and/or bromides are employed because of the high density which can be achieved, but other salts such as sulfates, nitrates, etc. can be employed. The only restriction is that the salts must be compatible with the particular thickening agent employed to thicken the aqueous fluid. By compatible it is meant, for example, that the salt does not detrimentally interfere with the thickening function of the thickening agent and/or undesirable quantities of precipitates are formed. As examples of useful water soluble salts, reference may be had to Table I, Column 3, of U. S. Patent No. 2,898,294.

One preferred aqueous wellbore service fluid contains a mixture of at least calcium bromide and zinc bromide to provide an aqueous solution having a density of at least about 15 ppg. The solution may also contain other water soluble salts such as calcium chloride and the like.

A preferred aqueous solution for use in deep wells requiring a fluid having a density greater than about 15 ppg is one which contains, as percent by weight:

ZnBr₂ about 5% to about 35%;

CaBr₂ about 25% to about 45%;

CaCl₂ about 5% to about 20%;

water about 30% to about 40%; and thickener about 0.5% to about 2%.

A preferred thickening agent for the above defined fluid having a density of above about 16.5 ppg is a tertiary amine of the formula $C_{18}H_{35}N(CH_2CH_2OH)_2$.

The thickening agent employed in the inven-20 tion comprises at least one of the thickening agents defined hereinbefore under Summary of the Invention. It is found that with certain solutions, a mixture of two or more thickeners may be preferred.

Preferably, X is an inorganic anion such as a sulfate, nitrate, perchlorate or halide. A halide, (Cl, Br or I) is preferred, Cl and Br being most preferred. X may also be an aromatic organic anion such as salicylate, naphthalene sulfonate, p and m chlorobenzoates, 3,5 and 3,4 and 2,4-dichlorobenzoates,

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t-butyl and ethyl phenate, 2,6 and 2,5-dichlorophenates, 2,4,5-trichlorophenate, 2,3,5,6-tetrachlorophenate, p-methyl phenate, m-chlorophenate, 3,5,6-trichloropicolinate, 4-amino-3,5,6-trichlorpicolinate, 2,4-dichlorophenoxyacetate, toluene sulfonate α,β-naphthols, p.p'bisphenol A. The thickening agent should be chosen such that the anion is compatible with the electrolyte present in the aqueous solution such that undesirable precipitates are not formed. Also, the specific anion chosen will depend to some degree on the specific amine structure.

The thickening agent is employed in an amount which is sufficient to increase the viscosity of the aqueous fluid at least 50 percent over the viscosity thereof without the addition of the thickener as measured on a Haake Rotovisco Viscometer at about 20°C and a shear rate of 160 sec⁻¹.

The exact quantity and specific thickener or mixture of thickeners to be employed will vary depending on the concentration of and specific soluble salt(s) employed to make up the solution, the viscosity desired, the temperature of use, the pH of the solution, and other similar factors. The concentration of the thickener can range from about 0.05 to about 5 percent, preferably from about 0.2 to about 3 percent of the aqueous wellbore service fluid. Simple laboratory procedures can be employed to determine the optimum conditions for any particular set of parameters. For example, when a non-protonated amine is employed as the thickener, the pH of the aqueous fluid can affect to some degree the effectiveness of particular amines. More acidic solutions are required for some amines to be dissolved therein. It is thought that this is

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because the amine must become protonated before it will become effectively dissolved in the fluid.

Specific wellbore service fluids found to be useful in the practice of the invention are set forth in the following Table I.

# 2			٠	118: -10-	5779				
oncentration Water Soluble Salt Percent by Weight ution #1 Solution									
Concentration Soluble Salercent by Wesolution #1 Solution	40	46.7	23	17.2	15.1 40.7 6.8	15.1	17.2	53	53
Water Soluble Salt	CaBr ₂	CaBr ₂	CaBr ₂	CaCl2 CaBr2	CaCl ₂ CaBr ₂ ZnBr ₂	CaCl ₂ CaBr ₂ ZnBr ₂	CaCl ₂ CaBr ₂	\mathtt{CaBr}_2	CaBr ₂
Concentration of Thickener Percent by wt.	1	0.5	. 0.5	0.5	0.5	0.38	0.75	1.08	1.5
Thickener	$c_{16}H_{33}$ N(CH ₃) ₃ salicylate	2	Oleyl Methyl bis(2-hydroxy-ethyl) ammonium chloride (OMB)	OMB	OMB	OMB and c_{16} H_{33} N(CH $_3$) $_3$ Cl	OMB	OMB	$c_{18}H_{37}N$ Br
Solution No.	-	7	ო	4	ν	9	7	89	σ

TABLE I (continued)

**		-11-			
tion Water e Salt by Weight Solution		6 28 33	6 28 33	6 28 33	3 8 6 3 8 6
Concentration Water Soluble Salt Percent by Weight Solution #1 Solution	53	10 34 20	10 34 20	10 34 20	10 34 20
Water Soluble Salt	CaBr ₂	CaCl CaBr ₂ ZnBr ₂	CaCl2 CaBr2 ZnBr2	CaCl CaBr ² ZnBr ²	CaCl CaBr2 ZnBr2
Concentration of Thickener Percent by wt.	1.5	*6.0	*6.0	* 6.0	*6.0
C Thickener P	снз $c_{18}^{\text{CH}_3}$, $c_{12}^{\text{CH}_2}$ CH ₂ CH ₂ CH ₂ OH) $c_{18}^{\text{CH}_3}$ Вг-	Bis(2-hydroxyethyl)oleylamine	Bis(2-hydroxyethyl)soyaamine	Bis(2-hydroxyethyl)tallowamine	Bis(2-hydroxyethyl)octadecyl- amine
Solution No.	10	11	12	13	14

a 75 percent active isopropanol solution of thickener in 35 ml of salt solution. of * 0.9 ml

-12-

Examples of other thickeners which can be employed include oleyl methyl bis(hydroxyethyl) ammonium chloride; octadecyl methyl bis(hydroxyethyl) ammonium bromide; octadecyl tris(hydroxyethyl) ammonium bromide; and octadecyldimethylhydroxyethyl ammonium bromide cetyldimethyl hydroxyethyl ammonium bromide; cetyl methyl bis(hydroxyethyl)ammonium salicylate; cetyl methyl bis(hydroxyethyl)ammonium 3,4-dichlorobenzoate; cetyl tris(hydroxyethyl)ammonium iodide; bis(hydroxyethyl) soyaamine; N-methyl, N-hydroxyethyl tallow 10 amine; bis(hydroxyethyl)octadecylamine; cosyl dimethylhydroxyethyl ammonium bromide; cosyl methyl bis(hydroxyethyl) ammonium chloride; cosyl tris(hydroxyethyl) ammonium bromide; docosyl dimethylhydroxyethyl ammonium bromide; docosyl methyl bis(hydroxyethyl)ammonium 15 chloride; docosyl tris(hydroxyethyl)ammonium bromide; hexadexyl ethyl bis(hydroxyethyl)ammonium chloride, hexadecyl isopropyl bis(hydroxyethyl)ammonium iodide; N,N-dihydroxypropyl hexadecylamine, N-methyl, N-hydroxyethyl hexadecylamine; N,N-dihydroxyethyl octadecylamine, 20 N, N-dihydroxypropyl oleylamine; N, N-dihydroxypropyl soya amine; N,N-dihydroxypropyl tallow amine; N-butyl hexadecyl amine; N-hydroxyethyl octadecylamine; N-hydroxyethyl cosylamine; cetylamine, N-octadecyl pyridinium chloride; N-soya-N-ethyl morpholinium ethosulfate; 25 methyl-1-oleyl amido ethyl-2-oleyl imidazolinium-methyl sulfate; methyl-1-tallow amido ethyl-2-tallow imidazolinium-methylsulfate.

It has been found that as the concentration of the soluble salt in the aqueous solution increases the thickener should be more hydrophilic. This can be achieved by employing thickeners having a specific combination of R_1 and R_2 - R_4 groups to provide such

-13-

hydrophillic character. It has also been found that the X^{-} component of the thickener affects, to some degree, the effectiveness of the thickener in specific aqueous solutions. For example, organic anions (X) generally are found to function more effectively in lower density fluids, e.g., less than 49% CaBr2, because of their solubility. Thickeners having an inorganic anion constituent are generally more effective over a broader density range than are thickeners containing an organic anion.

To prepare the aqueous wellbore service fluid of the present invention, the thickener is added to an aqueous solution to which has been dissolved a quantity of at least one water soluble salt to provide a solution having a desired density. Standard mixing procedures known in the art can be employed since heating of the solution and special agitation conditions are normally not necessary. Of course, if used under conditions of extreme cold such as found in Alaska, normal heating procedures should be employed. It has been found in some instances preferable to dissolve the thickener into a lower molecular weight alcohol prior to mixing it with the aqueous solution. The lower molecular weight alcohol (e.g., isopropanol) functions as an aid to solublize the thickener. Other such agents can 25 also be employed. A defoaming agent such as a polyglycol may be employed to prevent undesirable foam during the preparation of the service fluid.

In addition to the water soluble salts and thickening agents described hereinbefore, the aqueous 30 wellbore service fluid may contain other conventional constituents which perform specific desired functions,

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e.g., corrosion inhibitors, propping agents, fluid loss additives, and the like.

The fluids defined herein can be employed in standard wellbore treatment services employing techniques and equipment well known in the art. They may be used to control a well during certain wellbore operations such as during the perforation of liners and the like. They can also be employed as packer fluids, drilling fluids and the like.

The following examples are illustrative of aqueous wellbore service fluids of the present invention.

Example 1

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The rheological behavior of 0.5 percent oleyl methyl bis(2-hydroxyethyl) ammonium chloride in a 53% aqueous CaBr₂ wellbore service fluid over a shear rate range of 0.6-3900 sec⁻¹ was determined. The fluid was prepared by combining

(added as 0.29 g of commercially available 75% active Ethoquad 0/12) with 39.63 g of 53% ${\rm CaBr}_2$ aqueous solution.

The solution was prepared by adding the

25 Ethoquad 0/12 to the 53% CaBr₂ solution and shaking on
a mechanical shaker overnight at room temperature. A
clear, very viscoelastic solution with a layer of
stable foam on top resulted from this procedure.

The viscosity of the so prepared fluid was measured at three temperatures (approx. 23°C, 43°C and

60°C) as a function of shear rate. The lowest shear rate (0.66 sec-1) measurement was determined employing a Brookfield LTV viscometer with a UL adaptor. Twenty milliliters (ml) of solution were slowly removed from the bottom of a sample bottle to avoid introducing foam into the annulus between the rotating cylindrical bob and the stationary cup wall. The calculated viscosities at the shear rate of 0.6 rpm (0.66 sec⁻¹) are tabulated below in Table II for the three temperatures. readings changed with time so the viscosity in centipoise (cps) is reported in Table II as a range and not as a single value. This characteristic indicates the elastic, non-Newtonian nature of the solution.

TABLE II

15	Temperature °C	No. of Readings	Viscosity (cps)		
	23	4	470 - 620		
	43	3	720 - 750		
	59.5-60	3	370 - 530		

The viscosity of samples taken at higher shear rates were measured on a Haake Rotovisco using the NV double-gap cup system. The rotor is a hollow cylinder which fits over a stationary cylindrical stator on the inside with the other cylindrical stator being the inside wall of the stainless steel cup containing the sample solution. The eight ml of fluid required for the test were delivered from a hypodermic syringe. Torque is recorded on a single pen strip chart at successively increasing shear rates. Shear rate is increased stepwise by increasing the rotor rpm. 30

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Individual torque readings were taken at three temperatures. The temperature, shear rate and calculated viscosities are set forth in the following Table III.

5 <u>TABLE III</u>

	Temp.	Shear Rate (sec ⁻¹)	Viscosity (cps)
	25°C	5.4 10.8	54 81
	11	21.6	58.5
10	· · ·	43.1	45
10	11	86.2	32.6
	11	173	26.4
	· ·	345	20.3
	17	690	16.6
15	11	1380*	13.9
•	17	1380*	17.0
	17	2760	14.0
	19 .	173	26.4
	43.1	5.4	169
20	11	10.8	117
	17	21.6	99
	17	43.1	86.3
	1 1	86.2	60.8
0.5	11	173	37.1
25	 V	345	21.9
	;;	690	16.1
	11	1380.* 1380.*	12.2 13.4
	11	2760	11.4
30	11	173	38.5
	61.6	10.8	99
	17	21.6	67.5
	11	43.1	56.3
	и.	86.2	40.5
35	II.	173	28.4
	11 19	345	20.5
	11	690	14.5
	11	1380*	10.5
	"	1380*	12.0
40	11	2760	8.5
	11	173	28.7

Duplicate readings at different head scales of the instrument.

Example 2

A 40.3% aqueous $CaBr_2$ solution was thickened with 1% of cetyl trimethyl ammonium salicylate, [C_{16} - $H_{33}(CH_3)_3N^+$ salicylate as follows:

0.131 g of salicylic acid was mixed with 9.47 g of 0.1 M ${\rm C_{16}H_{33}N}^+({\rm CH_3})_3{\rm OH}^-$ solution and the resulting solution mixed with 30.4 g of a 53% aqueous CaBr₂ solution.

After dissolution, a clear, slightly yellow solution of high viscosity was formed.

Viscosity measurements were made on a Brook-field LTV viscometer with UL adaptor and the results are tabulated below:

TABLE IV

15	Shear Rate (sec ⁻¹)	Temp.	No. of Readings	Viscosity (cps)
	0.33	23	3	524 - 492
	0.65	23	Off Sc	ale
	0.66	27.5	3	317 - 307
20	0.66	37.5	3	145 - 144
	0.66	50	3	62 - 66

Example 3

A quaternary ammonium salt of the formula:

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$$C_{18}^{H_{35}}^{N^{+}}_{,} - (CH_{2}^{CH_{2}}OH)_{2} Cl^{-}$$
 (Ethoquad 0/12 75% active)

was employed to thicken an aqueous electrolyte solution as follows.

A solution was prepared by mixing 0.33 g
Ethoquad 0/12 with 49.67 g of a solution containing
17.2% CaCl₂/43.7% CaBr₂/39.1% H₂O, having a density of
about 15 ppg. After dissolving by mechanically
shaking, there was some foam. The aqueous solution
also contained 0.4 percent by weight of a corrosion
inhibitor comprising a mixture of N-octyl pyridinium
bromide and ammonium thiocyanate. Using the procedures of Example 1, the rheology of the fluid was
measured on the Haake Rotovisco NV system and the
calculated viscosity and shear rate data is set forth
in the following Table V.

TABLE V

	Тешр.	Shear Rate (sec-1)	Viscosity (cps)
	25.3°C	43.1	49.5
	•	86.2	47.3
5	•	173	44.7
	•	345*	40.8
	•	345*	41.8
	u	690	43.6
	•	1380*	38.5
10.	•	1380*	36.2
		2760	33.6
	•	3902*	32.3
	•	3902*	31.5
	•	1380	37. 4
15	W	690	44.6
	**	690	39.1
	•	345	41.1
	Ħ	173	44.4
	•	86.2	47.3
20	45.5	21.6	75.6
	*	43.1	72
		86.2	. 60.8
	*	173	50.1
		345	40.8
25	# #	345	36.5
	•	690	35.2
		1380	29.4
	*	2760*	26.1
		2760*	24.8
30	**	3902	23.1
	*	1380	30.3
		690	33.9
	*	345	40.9
		173	50.1
35	•	86.2	60.8
	62.7	10.8	126
	10	21.6	105.3
	11	21.6	117
	**	43.1	94.5
40	**	86.2	77.4
	10	173	63.5
	Ħ	345*	47.3
	17	345*	49.6
	W	690	35.7
45	18	1380	25.8

TABLE V (Continued)

	Temp.	Shear Rate (sec ⁻¹)	Viscosity (cps)
	62.7	2760	19.4
	u	3902*	18
5	u	3902*	17.3
	12	1380	26.1
	, ti	690	. 35.2
	13	345	46.9
	· • • •	173	6 4. 5
10	. w	86.2	82.1
	*	43.1	105.3
	85.7	21.6	75.6
	IJ	43.1	71.1
	· • • • •	86.2	61.4
15	. #	173	48.9
	. 17	345*	37.5
	Ħ	345*	39.1
	•	690	. 30
	W	1380	21.9
20	. u	2760	16.6
	11	3902	13.7
	•	1380	21.3
	. **	690 <i>-</i>	29.7
	•	348	37.9
25	t#	173	51.1
	•	86.3	64.4
	**	43.1	76.5

Duplicate readings at different head scales of the instrument.

30 Example 4

An aqueous solution was made up at room temperature containing 0.33 g of Ethoquad 0/12 (Example 1) in 49.67 g of a 15.5 ppg aqueous solution containing 15.1% CaCl₂, 40.7% CaBr₂, 6.8% ZnBr₂ and 37.4% water.

35 The resulting solution was clear and viscous. Viscosity measurements are set forth in the following Table VI.

A similar solution was prepared at room temperature as above except that a mixture of amines was employed. The solution contained 0.25 g of

-21-

Ethoquad 0/12, 0.13 g of $C_{16}H_{33}N^+(CH_3)_3$ Cl⁻ (Arquad 16-50/50% active) and 49.62 g of the 15.5 ppg density aqueous solution defined directly hereinbefore. The so prepared solution was viscous and clear. Viscosity measurements were made as described directly hereinbefore and are set forth in the following Table VII.

The solution containing the mixture of thickening agents demonstrated higher viscosities than did the solution containing the single thickener. This demonstrates the flexibility of being able to control the viscosity of high density aqueous fluids by the practice of the present invention.

TABLE VI

	_	_1.	
	Temp. °C	Shear Rate (sec ⁻¹)	Viscosity (cps)
	25	10.8	133
	W	21.6	113
5	11	43.1	93.9
	tr ·	86.2	80.8 68.5
	u	173 345*	44.4
	··· U	345*	47.6
10	n ·	690	29.7
10	, 11	1380	21.2
	u u	2760	16.6
	10 -	3902	15.0
	. U	1380	21.5
15	u	345	44.5
	**	173	75.7
	u v	21.6	111.9
	45	21.6	82.9
	11	43.1	74.3
20	U	86.2	63.8
	!!	173	54.2
	17	345*	40.3
	u	345*	40.5
25	11	690 1380	27.3 18.4
25	11	2760	12.9
	11	3902	11.4
	11	345	40.0
	65	43.1	26.6
30	"	86.2	27.4
30	11	173	27.4
	11	345	24.6
	11	690*	19.8
	1f	690*	20.0
35	11	1380	14.9
	11	2760	10.9
	11	3902	9.7
	87	173	4.4
	u	345	4.2
40	11	690	4.4
	11	1380	5.0
	11	2760	6.8
	11	3902	7.2
4.5	11	2760	6.5 5.6-5.1
45	••	1380	5.6-5.1

Duplicate readings at different head scales of the instrument.

-23-

TABLE VII

	Temp. °C	Shear Rate (sec-1)	Viscosity (cps)
	25	10.8	344
	17	21.6	260
5	16	43.1	164
	10	86.2	102
	U .	173	63.8
	W.	345*	42.1
	II .	345*	
10	II	690	48.7 37.8
	11	1380	
	11	2760	31.1
	H	3902	23.7-20.3
	11	1380	18.6-26.3
15	lf .	345	38.5
	11	173	75.7
	11	86	105
	11	86	173
		00	,153
	55.2	10.8	0.00
20	11	21.6	266
	tt	32.7	167
	11	86.2	128
	68	172*	101
	11	172*	79.2
25	11	345	86.5
	11	690	67.6
	lT .	1380	43.3
	1)	2760	40.5
	11	3902	34.7
30	ii		33.5
	11	1380	50.7
	11	345 173	62.2
		1/3	69.4
	89	10.8	
	11	21.6	117
35	11	32.7	78.3
	11		47
	11	86.2	31.3
	11	173	22
	11	3 4 5	12.5
40	ti .	690	7.8
	11	1380	5.5
	W.	2760	4.1
	11	3902*	4.3
	11	3902*	4.3
		690	4.3

^{45 *} Duplicate readings at different head scales of the instrument.

The following illustrates the fact that the amount of thickener and mixtures thereof which are effective can vary and that preliminary screening tests should be made prior to field use. The same thickening agent was employed as in the immediately preceding Example 4 except in different proportion: 0.17 g of Ethoquad 0/12 and 0.25 g Arquad 16-50 were mixed with 49.58 g of the same 15.5 ppg density solution defined in Example 4 hereinbefore. A very thick and viscoelastic fluid was formed. However, there was some insoluble materials floating on top. Upon heating to 70°C, the viscosity became low and the solution became unstable and formed two distinct phases. The fluid would not find general utility as a wellbore service fluid as contemplated herein.

Example 5

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A variety of different thickeners were screened to determine their effectiveness to thicken an aqueous solution containing 53% by weight of CaBr₂. The characteristics of the resulting fluids are tabulated in the following Tables VIII, IX and X. When viscosity data is shown, it was calculated employing data generated on a Haake Rotovisco NV system as described hereinbefore and set forth in one of the following Tables IX and X.

II	١
5	
TABLE	
7	

		_ , 1	100) (•			
Remarks	Clear solution. Doesn't crystallize	Clear solution but crystal- lizes on standing at room temperature	Cloudy solution	Clear solution	Clear solution	Opaque paste - crystallization	Clear solution	Clear solution
Viscosity	LOW	Some, not viscoelastic	Lox	Some, not viscoelastic	Less than at 50°C	Table IX	Table 1X	Table IX
O O	Roos	Roos	Room	45-50	70	. 30	09	06
Constituent 1.5% Wt	A. Chenant (CH3)3c1 (Afgdad 16-50)	B. CABH37N*(CH3)3C1 (AFquada 18-50)	C. C ₁₆ H ₃₃ +N Br		2	$D. C_{18}H_{37}^{\dagger}N$ Br	=	:

	Remarks Clear, but crystallizes	Clear solution	Clear solution
ontinued)	Viscosity Table X	Table X	Table X
TABLE VIII (Continued)	Temp. °C. Room	09	06
	Constituent 1.5% Wt E. Cl8H37N + CH2CH2OH)2Br	CH ₃	2

TABLE IX

	Temp. °C	Shear Rate (sec-1)*	Viscosity (cps)*
	30	5.4	492.6**
	11	10.8	349.2
5	17	21.6	219
	17	43.1	187.4
	11	86.2	300.1
	17	172.5	273.1
	19	345	136.6
10	10	689.9	106.9
	60	5.4	2728.9
	1f	21.6	1131.3
	17	86.2	533.3
	W	172.5	328.2
15	18	345	185.1
	W	689.9	129.3
	•	1379.8	73.7
	•	2759.7	39.9
	90	5.4	1278.2
20	18	10.8	1035
	W	21.6	794.5
	10	86.2	347.6
	N	172.5	211.6
	17	689.9	73.7
25	17	1379.8	44.7
	11	2759.7	23.3

^{*} Data as printed out from computer interfaced with Rotovisco NV.

^{**} Since thickener was not entirely in solution, the lower viscosities were expected.

TABLE X

	Temp. °C	Shear Rate (sec ⁻¹)*	Viscosity (cps)*
	30	5.4	1830.8
	W .	21.6	561.3
5	10	86.2	161.9
	12	172.5	86.4
	19	345	57.2
	18	1379.8	15.7
	60	5.4	1287.5
10	12	10.8	676.5
		43.1	215.5
		86.2	455.5
	10	172.5	281.7
	u .	689.9	95.8
15		1379.8	56.5
	90.3	5.4	1555.6
	10	10.8	1254.8
	11	21.6	872.2
	11	86.2	261.2
20	17	172.5	149
	n .	689.9	47.5
	u .	1379.8	27.9
	18	2759.7	17.1
	10	3902.4	17.1

25 * Data as printed out from computer interfaced with Rotovisco NV.

Example 6 and Comparative Tests

An electrolyte solution of about 16 ppg density was prepared containing 14% ZnBr₂; 37% CaBr₂; 12% CaCl₂ and 37% water. The fluid loss property of this fluid was determined employing a thickener as described hereinafter. A particulate fluid loss additive was also employed in some of the tests. The particulate fluid loss additive comprised a mixture of particulate aliphatic hydrocarbon resins. A comparative series of tests were run employing hydroxy ethyl cellulose as a fluid loss additive. The fluid loss tests

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were run on Brea Sandstone according to the API-RP39 standard fluid loss test using 1 inch by 1 inch Brea sandstone instead of filter paper. All the tests were conducted at 150°F and 600 psi. The thickener con-5 sisted of Ethoquad 0/12 (Example 1). To each 300 ml fluid sample containing the thickener, one drop of polypropylene oxide was added to control foaming. As a comparison, several solutions containing hydroxyethyl cellulose were tested in the same manner. The hydroxyethyl cellulose was a commercially available product 10 purchased under the trademark Vatrosol 250HHR. results of these tests are set forth in the following Table XII. This data illustrates the favorable fluid loss properties achieved by the practice of the invention.

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TABLE XII

	Time (Minutes)	Fluid Loss (milliliters)
	A. Test Solut	ion-1% Thickener
	1	2.6
5	4	6.4
	9	12.6
	16	21.2
	25	32.6
	30	39.0
	•	
10	B. Test Solut Fluid Loss	ion-1% Thickener and 0.5% Particulate Additive
	1	1.0
	4	1.6
	9	2.0
15	· 16	2.6
	25	3.0
	30	3.2
	C. Test Solut	ion (0.22%) Hydroxyethyl Cellulose
	1	4.6
20	4	14.0
	9	30.0
	16	57.0
	25	95.0
	30	Not measured

D.	Test So	lution-1.5	lb/gal	(0.22%)	Hydroxyethyl	Cellulose
	plus 5%	Particulate	Fluid	Loss A	dditive	

	plus_	5% Particulate Fluid Loss Additive	_
·	1	3.6	
	4	8.0	
5	9	11.0	
	16	12.2	
	25	13.0	
	30	13.2	

E. Test Solution-0.2% Thickener

10	1		2.4
	4		7.0
	9		15.6
	16		27.0
	25		47.0
15	30	•	59.0

F. Test Solution-0.2% Thickener plus 1.5% Particulate Fluid Loss Additive

_	Finid ross vddifine		
	1	1.0	
	4	7.0	
20	9	3.2	
	16	4.8	
	25	7.2	
	30	8.4	

Example 7 and Comparative Tests

Several thickening agents were evaluated for use in aqueous fluids having densities of greater than 15 ppg. In each example (data set forth in the following Table XII), 0.9 ml of a thickener was added to 35 ml of the indicated high density fluid.

The thickener was employed as a 75 percent active solution in isopropanol. The resulting mixtures

were shaken on a mechanical shaker for 3 hours and three days later the viscosities were measured at room temperature (about 75°F) employing a Brookfield viscometer at a shear rate of 60 rpm employing a number 2 or 4 spindle. The data employing the number 4 spindle is marked with an asterisk. The data indicates that the thickening agents were more effective in the higher density fluids.

The results are tabulated in the following 10 Table XII.

TABLE XII

		Viscosity (cps)		
	Thickening Agent	15.5 ppg (2)	16.5 ppg (3)	17.5 ppg (4)
15	<pre>bis(2-hydroxyethyl)- oleylamine</pre>	45	3250*	850*
	<pre>bis(2-hydroxyethyl)- soyaamine</pre>	, 40	2500*	550*
20	<pre>bis(2-hydroxyethyl)- tallowamine</pre>	40	1900*	420
	<pre>bis(2-hydroxyethyl)- octadecylamine</pre>	45	300*	60
	none	28	25	25
	none	50*	40*	40*

- 25 (2) Solution contained 15% CaCl $_2$; 41% CaBr $_2$; 6% ZnBr $_2$ and 38% H $_2$ O.
 - (3) Solution contained 10% CaCl₂; 34% CaBr₂; 20% ZnBr₂ and 36% H₂O.
- (4) Solution contained 6% CaCl $_2$; 28% CaBr $_2$; 33% ZnBr $_2$ and 33% H $_2$ O.

Example 8

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A solution was prepared by dilution of a 40 g sample of 1.5% solution (0.6 g of $C_{22}H_{46}(CH_3)_2N^9C_2H_4OHBr^9$ + 39.4 g of 17.5 ppg fluid) with 80 g of 17.5 ppg fluid to provide 0.5% of the surfactant in the solution. The 17.5 ppg fluid contained 6% CaCl; 28% CaBr₂; 33% ZnBr₂, and the remainder water. The sample was heated to approximately 85°C with occasional shaking to produce a clear, homogeneous solution which was hazy at room temperature. The viscosity was measured in a Haake Rotovisco as described in Example 1. These data are tabulated in Table XIII and show good thickening at room and intermediate temperatures, and filling off at 85°C. The viscosities are from a computer printout interface with the Haake Rotovisco.

-34-

TABLE XIII

	Temp. °C	Shear Rate (sec-1)	Viscosity (cps)
	26	5.4	1025
	•	10.8	789.7
5	•	21.6	439.2
	•	43.1	241.7
	•	86.2	149.7
	. •	172.5	107.1
	•	345	67.6
10	•	689.9	48.9
	•	1379.8	38.7
	•	2759.7	31.8
	•	3902.4	29.6
	54	. 5.4	635
15	•	10.8	436.7
	•	21.6	317.5
	•	43.1	405.3
	•	86.2	166.6
		172.5	64.9
20	•	345	35.7
	•	689.9	25.1
		1379.8	17
	•	2759.7	14.1
	•	3902.4	13.2
25	855	86.2	10.1
	•	172.5	9.3
	•	345	8.7
	•	689.9	8.4
	•	1379.8	8.4
30	•	2759.7	7.1
	#	2759.7	9.7
	•	3902.4	10.4

-35-

Example 9

A solution was prepared by diluting a 50 g sample of 1.5% solution (0.75 g of $C_{20}H_{41}(CH_3)_2N^{\theta}C_2H_4OHBr^{\theta}$ + 49.25 g of 19.2 ppg fluid) with 50 g of 19.2 ppg fluid to provide 0.75% of the thickening agent in the fluid. The fluid contained 56% ZnBr2, 19% CaBr2 and 25% H₂O. The sample was heated to 85°C to provide rapid dissolution. The appearance at this temperature was a clear viscous solution which was also clear when cooled to room temperature (21-22°C). After standing a number of days, the sample tended to become hazy although warming (approximately 30-35°C) restored the original clear appearance. The viscosity of the fluid was determined as a function of shear rate on a Haake Rotovisco with a computer printout as previously described. The results are set forth in the following Table XIV.

-36-

TABLE XIV

	Temp. °C	Shear Rate (sec ⁻¹)	Viscosity (cps)
	27	10.8	104.8
_	11	21.6	116.8
5	ti	43.1	87.8
	11	86.2	66.5
	"	172.5	50.1
	**	345	39.4
	. 11	689.9*	34.3
10	11	689.9*	34.3
	12	1379.8	29.5
		2759.7	25.8
	11	3902.4	24.5
	55 -	5.4	138.6
15	11	10.8	125.7
	v	5.4	138.6
	17	10.8	125.7
	t7	21.6	123.3
	u	43.1	117.2
20	. 10	86.2	99.3
		172.5	79.1
	tf	345	63.8
	₩	689.9	46.5
	st	1379.8	35.7
25	11	2759.7	27.6
	W	3902.4	24.6
	85	21.6	50.8
	tf	43.1	47.1
	17	86.2	44.3
30	11	172.5	42.6
	t)	345	39.1
	tt	689.9*	36.8
	tī	689.9*	36.8
	11	1379.8	31
35	11	2759.7	25.6
	11	3902.4	22.9

Duplicate readings at different heat scale of instrument.

THE EMBODIMENTS OF THE INVENTION IN WHICH AN EXCLUSIVE PROPERTY OR PRIVILEGE IS CLAIMED ARE DEFINED AS FOLLOWS:

l. An aqueous wellbore service fluid comprising: water, a sufficient quantity of at least one water soluble salt to increase the density of said fluid to within a range of from 12 to 21 lbs/gal, and a sufficient quantity of at least one thickener soluble in said fluid to increase the viscosity of said fluid to at least 50% over the viscosity of the salt containing fluid, said thickener being at least on member selected from the group consisting of: (a) an amine corresponding to the formula

$$R_1 - N$$
 wherein R_3

 R_1 is at least about a C_{16} aliphatic group which may be branched or straight chained and which may be saturated or unsaturated;

 $\rm R_2$ and $\rm R_3$ are each independently, hydrogen or a $\rm C_1$ to about $\rm C_6$ aliphatic group which can be branched or straight chained, saturated or unsaturated and which may be substituted with a group which renders the $\rm R_2$ and/or $\rm R_3$ group more hydrophilic; (b) salts of said amine corresponding to the formula

$$R_1 - N^+ - H \quad X^-$$
 wherein R_3

 R_1 , R_2 and R_3 are the same as defined hereinbefore and X is an inorganic or organic salt forming anion; or (c) a quaternary ammonium salt of said amine corresponding to the formula

$$R_1 - N^+ - R_4 \quad X^-$$
 wherein R_3

 R_1 , R_2 , R_3 and X^- are the same as hereinbefore defined and R_4 independently constitutes a group which has previously been set forth for R_2 and R_3 , none of R_1 , R_2 , R_3 or R_4 are hydrogen, and the R_2 , R_3 and R_4 groups of the amine salt and quaternary ammonium salt may be formed into a heterocyclic 5 or 6 member ring structure which includes the nitrogen atom of the amine.

- 2. The wellbore service fluid of Claim 1 wherein the thickening agent is employed in an amount ranging from about 0.05 to about 5 percent by weight of the fluid.
- 3. The wellbore service fluid of Claim 1 wherein the water soluble salt comprises a compination of calcium chloride, calcium bromide and zinc bromide in an amount and weight ratio to provide a density of at least 15 pounds per gallon of fluid.

- 4. The wellbore service fluid of Claim 1 wherein the density of the fluid is at least about 15 pounds per gallon of fluid.
- 5. The wellbore service fluid of Claim 1 wherein the thickening agent comprises at least one member selected from $C_{16}^{H}_{33}^{N}(CH_{3})_{3}$ salicylate; oleyl methyl bis(2-hydroxyethyl)ammonium chloride;

$$c_{16}^{H_{33}N^{+}(CH_{3})_{3}CL^{-}}; c_{18}^{H_{37}} + N \longrightarrow Br^{-}; c_{18}^{H_{37}N^{+}(CH_{2}CH_{2}OH)_{2}Br^{-}}; c_{18}^{H_{37}N^{+}(CH_{3}CH_{2}OH)_{2}Br^{-}};$$

bis(2-hydroxyethyl)oleylamine; bis(2-hydroxyethyl)soya-amine; bis(2-hydroxyethyl)tallowamine; bis(2-hydroxyethyl)-octadecylamine; $C_{18}^{H_{35}N(CH_{2}CH_{2}OH)_{2}}$; $C_{22}^{H_{45}(CH_{3})_{2}N^{+}CH_{2}CH_{2}OHBr^{-}}$ or $C_{20}^{H_{41}(CH_{3})_{2}N^{+}CH_{2}CH_{2}OHBr^{-}}$.

- 6. The wellbore service fluid of Claim 1, wherein the water soluble salt comprises from 5% to 35% ZnBr₂; from 25% to 45% CaBr₂; from 5% to 20% CaCl₂ from 30 to 40% water, and from 0.5 to 20% of said thickening agent, all percentages being in percent by weight of the fluid.
- 7. The wellbore service fluid of Claim 1 wherein X is Cl or Br.
- 8. In the method of perforating an interval of casing in a wellbore wherein a sufficient quantity of fluid is placed in the wellbore adjacent to the interval to be perforated to maintain pressure on the formation which is at least as great as the formation pressure, the improvement which comprises as the fluid the aqueous wellbore service fluid of Claim 1, 5 or 6.

9. In the method of servicing a wellbore wherein an aqueous fluid is placed in the wellbore and in contact with an oil and/or gas producing formation the improvement which comprises employing as the fluid the aqueous wellbore service fluid of Claim 1, 5 or 6.

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PATENT AGENTS

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